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(54) Title: METHODS AND DEVICES FOR TISSUE REPAIR

(57) Abstract: Methods for treating diseased or damaged tissue in a subject are disclosed, involving administering to said subject at a site wherein diseased or damaged tissue occurs, cells of a type(s) normally found in healthy tissue corresponding to the diseased or damaged tissue, and/or suitable progenitor cells thereof, in association with bioresorbable beads or particles and optionally a gel and/or gel-forming substance. Where the cells and/or suitable progenitor cells thereof are chondrocytes, embryonic stem cells and/or bone marrow stromal cells, the methods of the invention are suitable for treating, for example, articular cartilage degeneration associated with primary osteoarthritis. Also disclosed is a device having tissue-like characteristics for treating diseased or damaged tissue in a subject, wherein the device comprises cells of a type(s) normally found in healthy tissue corresponding to the diseased or damaged tissue, and/or suitable progenitor cells thereof, in association with bioresorbable beads or particles and optionally a gel and/or gel-forming substance.



WO 02/062357 A1

METHODS AND DEVICES FOR TISSUE REPAIR

Field of the Invention:

The present invention relates to methods and devices for treating diseased or
5 damaged tissue, particularly articular cartilage degeneration associated with primary
osteoarthritis, and other articular cartilage damage caused by, for example, sporting
injuries or trauma. The present invention may also be applied to tissue augmentation
(e.g. for cosmetic reasons).

Background of the Invention:

Articular cartilage is found lining the bones within bone joints (e.g. the knee)
where it allows for stable movement with low friction and provides resistance to
compression and load distribution. The articular cartilage appears as a simple,
avascular matrix of hyaline cartilage but, in fact, consists of a relatively complex
15 formation of chondrocytes and extracellular matrix (ECM) organised into four zones
(i.e. the superficial, transitional, middle and calcified zones) based upon matrix
morphology and biochemistry. In turn, each of these zones consists of three distinct
regions (i.e. the pericellular, territorial, and interterritorial regions). Chondrocytes,
which comprise less than 5% of the volume of human articular cartilage, replace
20 degraded ECM molecules and are thereby essential for maintaining tissue integrity (i.e.
size and mechanical properties). The ECM includes a number of components including
collagen (primarily, Type II collagen), glycoproteins, proteoglycans and tissue fluid
which comprises up to about 80% of tissue weight of articular cartilage. The collagen
component provides a fibre mesh structure to the ECM and the glycoproteins are
25 thought to assist in the stability of the structure. The proteoglycans comprise large
aggregating monomers (i.e. aggregans) which fill the inter-fibre spaces and, because of
their ability to attract water, are believed to account for much of the resiliency and load
distribution properties of articular cartilage. Finally, the tissue fluid, which includes a
source of nutrients and oxygen, provides the articular cartilage with the ability to resist
30 compression and return to its regular shape following deformation (for a review, see
Temenoff and Mikos, 2000).

Joint pain resulting from articular cartilage degeneration or injury is a common
condition which afflicts people of all ages. Its major causes are primary osteoarthritis
and trauma causing loss of cartilage (Buckwalter and Mankin, 1998). Recently, it has
35 been estimated that up to 43 million people in the United States of America alone suffer

from some form of arthritis (see "Arthritis Brochure" at <http://orthoinfo.aaos.org/>), while cartilage damage arising from sporting injuries is also prevalent.

Unfortunately, and owing in part to its complex structure (Temenoff and Mikos, *supra*), articular cartilage has extremely little ability for self repair and, as a
5 consequence, articular cartilage degeneration and injuries persist for many years and often lead to further degeneration (i.e. secondary osteoarthritis).

Treatment options for articular cartilage degeneration can be grouped according to four principles, i.e. replacement, relief, resection and restoration. Replacement of articular cartilage involves the use of a prosthesis or allograft. Relief of symptoms can
10 be achieved by an osteotomy operation, which removes a portion of one of the bones in the defective joint so as to decrease loading and stress. Resection refers to surgical removal of the degenerated articular cartilage and subsequent uniting of the healthy, surrounding articular cartilage tissue. Such resection operations may or may not involve the use of interposition arthroplasty. Lastly, restoration refers to healing or
15 regeneration of the joint surface, including the articular cartilage and the subchondral bone. This may involve an attempt to enhance self repair (e.g. through use of pharmaceutical agents such as growth factors, or subchondral drilling, abrasion or microfracture to "recruit" pluripotent stem cells from the bone marrow), or otherwise, regenerating a new joint surface by transplanting chondrocytes or other cells having the
20 ability to regenerate articular cartilage.

Considerable research has been conducted in recent years on the development of suitable "restoration" treatments or, more specifically, treatments involving regeneration of a new joint surface (sometimes referred to as "biological resurfacing"). Such treatments may be less traumatic to a patient than an osteotomy or prosthetic
25 replacement, and offer advantages over the use of allografts which may not be immunologically tolerated and which may contain foreign pathogens, or multiple autografts which, inevitably, cause damage at another site on the patient. One "biological resurfacing" treatment that has been proposed involves the harvesting of chondrocytes from an articular cartilage biopsy from the patient (Freed et al., 1999).
30 These cells are expanded in culture, and administered back to the patient by injection under a periosteal flap, which is sutured to ensure that the expanded chondrocytes remain at the site requiring repair. While this treatment has shown considerable promise in human trials over the past decade (Temenoff and Mikos, *supra*), the need for a periosteal flap adds an additional restriction to the technique, and the act of
35 sewing the periosteal flap over the injected chondrocytes can lead to damage to the adjacent tissue. Additionally, there is no evidence to suggest that the expanded cells

remain phenotypically and functionally as chondrocytes; indeed, they may have de-differentiated into fibroblast-like cells that produce mechanically inferior tissue.

A potential alternative to the use of the above system of autologous cells and periosteal flap, is the use of preformed porous scaffolds that approximates the desired
5 shape and form of the diseased or damaged tissue, and which have been seeded with chondrocytes and cultured for at least 2 to 3 weeks. The tissue equivalent that forms is then implanted at the required site (Thomson et al., 1995). Recent work with collagen-based scaffolds has been promising, however most of the current research being conducted in this area is concerned with identifying suitable synthetic polymer
10 materials for scaffolds, since these may be produced in large amounts and should overcome the concerns surrounding the possibility of incomplete pathogen removal from donor collagen (Temenoff and Mikos *supra*). Particular examples of synthetic polymer materials being researched are fibres of FDA-approved polymers, poly(glycolide) (PGA), poly(lactide) (PLA) and copolymers poly(lactide-co-glycolide)
15 (PLGA). These polymer fibres, which may be woven into a mesh, are biodegradable and therefore offer advantages over non-degradable polymers in that their gradual degradation steadily creates room for tissue growth and, secondly, they eliminate the need for surgical removal of the scaffold following restoration of the articular cartilage.

The use of scaffolds does, however, have the substantial disadvantage of
20 necessitating surgery for implantation. Accordingly, other research groups have directed their efforts towards the development of polymers, which may be injected with chondrocytes and, subsequently, become cross-linked *in situ* to form a scaffold matrix. For example, fibrinogen and thrombin can be combined and injected wherein a degradable fibrin mesh is formed (Sims et al., 1998), and alginate has also been
25 investigated since this may be cross-linked with calcium (Rodriguez and Vacanti, 1998). Alginate has, however, been found to be immunogenic (Kulseng et al., 1999) and invokes a greater inflammatory response than synthetic polymer materials (Cao et al., 1998). Thus, research has also been conducted with injectable synthetic polymer gel materials including copolymers of ethylene oxide and propylene oxide PEO-co-
30 PPO (Cao et al., *supra*) and photopolymerizable end-capped block copolymers of poly(ethylene oxide) and an α -hydroxy acid (Hubbell, 1998).

The present invention relates to an alternative method for tissue regeneration, particularly articular cartilage regeneration, wherein chondrocytes and/or other suitable progenitor cells are bound to, or otherwise blended with, bioresorbable beads or
35 particles for administration to a subject at a site where tissue regeneration is required. It is believed that the method avails itself of many of the advantages of biodegradable

polymer scaffolds discussed above, including the ability to be administered by injection if desired. Additionally, and while not wishing to be bound by theory, it is thought that the use of beads or particles may provide mechanical and space-filling benefits while tissue regeneration is progressing by offering physical support and resistance to
5 compression.

Disclosure of the Invention:

Thus, in a first aspect, the present invention provides a method for treating diseased or damaged tissue in a subject, said method comprising administering to said
10 subject at a site wherein said diseased or damaged tissue occurs, cells of a type(s) normally found in healthy tissue corresponding to said diseased or damaged tissue, and/or suitable progenitor cells thereof, in association with bioresorbable beads or particles and, optionally, a gel and/or gel-forming substance.

The said cells and/or progenitor cells may be associated with the beads or
15 particles simply through mixing and may therefore not necessarily be bound to the beads or particles. The cells and/or progenitor cells may be mixed with the beads or particles by low shear agitation in a suitable vessel. The gel and/or gel-forming substance may be simultaneously mixed with the cells and/or progenitor cells and beads or particles, or alternatively mixed subsequently. However, preferably, the cells
20 and/or progenitor cells are associated with the beads or particles by being bound thereto. This may be achieved by expanding the cells and/or progenitor cells in the presence of the beads or particles.

Thus, in a second aspect, the present invention provides a method for treating diseased or damaged tissue in a subject, said method comprising the steps of;
25 (i) obtaining cells of a type(s) normally found in healthy tissue corresponding to said diseased or damaged tissue and/or suitable progenitor cells thereof,
(ii) expanding said cells and/or progenitor cells in the presence of bioresorbable beads or particles whereby said expanded cells and/or progenitor cells become bound to the said beads or particles, and
30 (iii) administering to said subject the beads or particles with said cells and/or progenitor cells bound thereto, optionally in a gel and/or gel-forming substance, at a site wherein said diseased or damaged tissue occurs.

It will be appreciated by persons skilled in the art that between steps (i) and (ii) above, an additional expansion step(s) may be carried out. Such additional expansion
35 step(s) may involve growth of the cells in, for example, monolayer(s).

It will also be appreciated by persons skilled in the art that it is not necessary to expand the cells and/or progenitor cells in the presence of the beads or particles at all and that, alternatively, the cells and/or progenitor cells could be expanded and, subsequently, bound to the beads or particles.

- 5 Thus, in a third aspect, the present invention provides a method for the treatment of diseased or damaged tissue in a subject, said method comprising the steps of;
- (i) obtaining cells of a type(s) normally found in healthy tissue corresponding to said diseased or damaged tissue and/or suitable progenitor cells thereof,
 - (ii) expanding said cells and/or progenitor cells,
 - 10 (iii) binding said expanded cells and/or progenitor cells to bioresorbable beads or particles, and
 - (iv) administering to said subject the beads or particles with said cells and/or progenitor cells bound thereto, optionally in a gel and/or gel-forming substance, at a site wherein said diseased or damaged tissue occurs.

- 15 The said cells and/or progenitor cells are selected such that they are of a type(s) suitable for regeneration of the particular diseased or damaged tissue type (e.g. mature differentiated cells of the tissue type to be treated). Thus, by way of example, for the treatment of diseased or damaged skin, the cells used in the methods of the present invention shall be fibroblasts and/or progenitor cells thereof. Where the tissue to be
- 20 regenerated is bone, the cells shall be osteoblasts and/or progenitor cells thereof, while for the treatment of fatty tissues, the cells shall be adipocytes and/or progenitor cells thereof.

- Preferably, the methods of the present invention are used for treating (e.g. repairing) articular cartilage degeneration or injury. In this regard, articular cartilage
- 25 tissue regeneration may be achieved at the site of articular cartilage degeneration or injury, and the bioresorbable beads or particles are gradually degraded so that removal of the beads or particles following regeneration is not required. In this application of the methods of the present invention, the cells used are chondrocytes and/or progenitor cells thereof. Further, as mentioned above, it is thought that while tissue regeneration
- 30 is progressing, the beads or particles provide mechanical and space-filling benefits. That is, they may provide a load-bearing cushion to the articular cartilage degeneration or injury by offering physical support to the bone joint, reduced friction during joint movement and resistance to compression. In addition, where the beads or particles are administered in a gel or gel-forming substance, the beads or particles appear to prevent
- 35 gel contraction, which might otherwise adversely affect space-filling of the tissue defect.

The chondrocytes and/or progenitor cells may be harvested by any of the methods common to the art, but most conveniently, by tissue biopsy. Suitable chondrocyte progenitor cells are undifferentiated cells such as embryonic stem cells and bone marrow stromal cells. Preferably, the chondrocytes and/or progenitor cells
5 are obtained from the subject to be treated.

The expansion step in the methods of the second and third aspects, preferably expand the cells and/or progenitor cells 5 to 2000-fold, more preferably, 10 to 100-fold, by any of the methods common to the art. For example, expansion may be achieved by cell culture in a suitable dish (such as a petri dish, with or without, for example, an agar
10 gel being present), but more preferably, is conducted in a bioreactor where the culture medium is agitated and aerated. The expansion may, however, involve more than one stage. For example, chondrocytes and/or progenitor cells thereof may first be grown as a monolayer in a suitable dish, wherein cell spreading may be mediated by serum adhesion proteins such as fibronectin (Fn) and vitronectin (Vn), and subsequently
15 grown in a bioreactor. As mentioned above, the expansion, or a portion of the expansion, may or may not be conducted in the presence of bioresorbable beads or particles. Also, when beads or particles are present during the expansion, or a portion of the expansion, the cells and/or progenitor cells may be removed and "re-seeded" onto bioresorbable beads or particles. In this case, the first mentioned beads or
20 particles may not necessarily be bioresorbable beads or particles. Where the expansion involves culturing in a bioreactor, it is convenient to add bioresorbable beads or particles to the culture medium. However, where the expansion is conducted without beads or particles, it is necessary, as is clear from the above, to subsequently bind the expanded cells and/or progenitor cells to bioresorbable beads or particles.

25 A simple bioreactor that is suitable for expansion of cells (e.g. chondrocytes) and/or progenitor cells for use in the methods of second and third aspects, is a spinner flask. Alternatively, expansion of the cells and/or progenitor cells may be achieved with a tumbler-type bioreactor (eg: Synthecon™ Inc. STLV™ Rotary Cell Culture System) which may or may not be equipped with internal vanes to assist in movement
30 of the cells, culture medium and bioresorbable beads or particles, if present.

Where chondrocytes are used, culturing in a spinner flask or tumbler-type bioreactor should ensure maintenance of cell phenotype. However, where the expansion involves culturing in an essentially still culture medium, it may be necessary to take steps to prevent de-differentiation of the chondrocytes. In both cases, the
35 culture medium may include supplements, such as ascorbate or growth factors, which control the cell growth and characteristics.

The bioresorbable beads or particles utilised in the methods of the present invention are preferably sized such that they are readily injectable. Accordingly, the bioresorbable beads or particles preferably have a diameter or dimensions sized in the range of about 20 to 2500 μm , more preferably, with an average size of about 50 to 200 μm . Suitable bioresorbable beads may be of a regular shape (e.g. spheroid such as microspheres, ovoid, disc-like or rod-like) or a mixture of regular shapes. On the other hand, suitable bioresorbable particles will generally be comprised of a large variety of irregular shaped particles as would typically be produced from crushing or pulverising solid substances.

10 The bioresorbable beads or particles may be comprised of any pharmaceutically acceptable polymer including biologically-based polymers such as gelatin and collagen (especially type I and/or type II), and synthetic polymers such as those, which have been used in, cell scaffolds (i.e. PGA, PLA and PLGA), and mixtures of biologically-based and synthetic polymers. Alternatively, the bioresorbable beads or particles may
15 be comprised of other pharmaceutically acceptable non-polymeric substances including bone particles (e.g. crushed bone and particles of demineralised bone). Also, the bioresorbable beads or particles may be comprised of a mixture of such polymers and non-polymeric substances.

Preferably, the bioresorbable beads or particles are of a size and density that
20 allows thorough movement of the beads or particles in a spinner flask or tumbler-type bioreactor. This may assist in cell expansion and, where chondrocytes are being used, maintenance of chondrocyte phenotype.

The bioresorbable beads or particles may be functionalised or coated in a suitable material to enhance cell adherence (e.g. an antibody or fragment thereof which
25 binds to a cell-surface antigen, or ECM proteins such as collagen Type I, II, VI, IX, XI, etc.) and/or, where chondrocytes are being used, may also be coated with an agent to assist in the maintenance of phenotype (e.g. a type II collagen). Additionally, the beads or particles may comprise other beneficial agents such as growth factors (e.g. TGF β , EGF, FGF, IGF-1 and OP-1, etc.), glycosaminoglycans (GAGs) (e.g. aggrecan,
30 decorin, biglycan, fibromodulin) and hydrophilic compounds (e.g. polylysine, chitosan, hyaluronan).

Preferably, the beads or particles, with suitable cells and/or progenitor cells associated therewith, are administered to a subject in a gel and/or gel-forming substance. However, additionally or alternatively, the beads or particles with suitable
35 cells and/or progenitor cells associated therewith, may be administered in combination

with a suitable pharmaceutically acceptable carrier (e.g. physiological saline, sterile tissue culture medium, etc.).

Suitable gel and/or gel-forming substances are preferably bioresorbable and of a type that ensures that the beads or particles are substantially retained at the site of administration. The gel and/or gel-forming substance may, therefore, comprise an adhesive material(s) (e.g. fibrin and/or collagen, or a transglutaminase system) to adhere the gel or formed gel to the tissues surrounding the site of administration. Alternatively, or additionally, the beads or particles may be substantially retained at the site of administration by entrapping the gel and/or gel-forming substance containing the beads or particles within tissue (e.g. the dermal and/or adipose tissue(s)) or under a tissue (e.g. a periosteal flap) or other membranous flap (e.g. a collagen membrane).

Suitable gels and gel-forming substances may comprise a biologically-based polymer (i.e. a natural or treated natural polymer) such as a collagen solution or fibrous suspension, hyaluronan or chitosan (hydrolysed chitin), or a synthetic polymer such as a photopolymerizable end-capped block copolymer of poly(ethylene oxide) and an α -hydroxy acid. The gel and/or gel-forming substance may also comprise other beneficial agents such as growth factors (including those mentioned above), glycosaminoglycans (GAGs) and hydrophilic compounds (such as those mentioned above).

In the methods of the second and third aspects, the cells and/or progenitor cells bound to the beads or particles, when ready for administration, may be confluent or sub-confluent. An average between about 3 and 500 cells and/or progenitor cells are preferably associated with each bioresorbable beads or particles. The numbers will, however, vary depending upon the characteristics (e.g. composition and size) of the beads or particles. For administration, it is preferred to use 1×10^5 to 1×10^9 cells and/or progenitor cells bound per 1 cm^3 of beads or particles.

Where chondrocytes are used, the chondrocytes bound to the beads or particles may be administered to the subject, before or after the chondrocytes have commenced secreting extracellular matrix. The latter is, however, less preferred since the extracellular matrix can lead to the formation of aggregates, which may not be readily injectable.

In the method of the third aspect of the present invention, the cells and/or progenitor cells are first expanded and then (i.e. subsequently), bound to bioresorbable beads or particles. This may be achieved in a suitable dish (e.g. a petri dish) or in tissue culture flasks. Again, the bioresorbable beads or particles may be functionalised or coated in a suitable material to enhance cell adherence, and/or coated with an agent to assist in the maintenance of chondrocyte phenotype. The beads or particles may also

comprise other beneficial agents such as growth factors, glycosaminoglycans (GAGs) and hydrophilic compounds.

In the method of the third aspect of the invention, the beads or particles with bound cells and/or progenitor cells can be administered to the patient immediately after
5 step (iii), or after further culturing of the cells and/or progenitor cells on the beads or particles.

The administration of the cells and/or progenitor cells in association with the beads or particles and gel and/or gel-forming substance is preferably by injection or arthroscopic delivery.

10 The methods of the present invention are primarily intended for human use, particularly in relation to treatment of articular cartilage tissue degeneration or injury (e.g. in the knee, fingers, hip or other joints). However, it is also anticipated that the methods may well be suitable for veterinary applications (e.g. in the treatment of articular cartilage degeneration or injury in race horses, and in the treatment of articular
15 cartilage degeneration or injury in companion animals).

The present invention also contemplates the production of a tissue-like device that may be surgically implanted into a subject for the treatment of diseased or damaged tissue.

Thus, in a fourth aspect, the present invention provides a device having tissue-
20 like characteristics for treating diseased or damaged tissue in a subject, wherein said device comprises cells of a type(s) normally found in healthy tissue corresponding to said diseased or damaged tissue, and/or suitable progenitor cells thereof, in association with bioresorbable beads or particles and optionally a gel and/or gel-forming substance.

The device may be prepared by culturing said cells and/or progenitor cells in
25 association with bioresorbable beads or particles and optionally a gel and/or gel-forming substance, for a period of time sufficient so as to form a tissue-like mass. The cells and/or progenitor cells may or may not be bound to the bioresorbable beads or particles. The bioresorbable beads may have fully degraded prior to implantation of the device, but preferably, the beads or particles are substantially intact within the device at
30 the time of implantation.

In a fifth aspect, the present invention provides a method for treating diseased or damaged tissue in a subject, said method comprising implanting into said subject at a site wherein said diseased or damaged tissue occurs, a device according to the fourth aspect.

It will be readily appreciated by persons skilled in the art that a combination of different types of cells, potentially on the same or different types of beads, could be used to effect repair of the diseased or damaged tissue.

It will also be readily appreciated by persons skilled in the art that the present invention may be applied to tissue augmentation (e.g. treatment of scars or facial wrinkles).

By the term "bound" we refer to any mechanism by which cells and/or progenitor cells may adhere to a bioresorbable bead or particle so that substantially all of said cells and/or progenitor cells bound to a particular bioresorbable bead or particle remain bound to that bead or particle. Such mechanisms include binding of chondrocytes and/or progenitor cells to said bead via an antibody (which may be covalently bound to the bead), or via an ECM protein (eg. collagen Type I, II, VI, IX, XI, etc.), or fragments thereof, which may also be covalently bound to the bead.

By the term "gel" we refer to any viscous or semi-solid solution or suspension which is capable of retarding settling of bioresorbable beads or particles as described above (c.f. bioresorbable beads or particles will readily settle out of physiological saline). Such solutions and suspensions preferably do not flow through a #2 Zahn Cup (Gardco, Inc.) (44 ml placed in the #2 Zahn Cup) at 37°C and atmospheric pressure in less than 30 seconds. More preferably, such solutions or suspensions do not flow through a #4 Zahn Cup (Gardco, Inc.), that is less than 5% of the initial volume (44 ml placed in the #4 Zahn Cup) flows through after 2 minutes at 37°C and atmospheric pressure.

The terms "comprise", "comprises" and "comprising" as used throughout the specification are intended to refer to the inclusion of a stated step, component or feature or group of steps, components or features with or without the inclusion of a further step, component or feature or group of steps, components or features.

Any discussion of documents, acts, materials, devices, articles or the like which has been included in the present specification is solely for the purpose of providing a context for the present invention. It is not to be taken as an admission that any or all of these matters form part of the prior art base or were common general knowledge in the field relevant to the present invention as it existed in Australia or elsewhere before the priority date of each claim of this application.

The invention is hereinafter further described by way of the following non-limiting examples and accompanying figures.

Brief description of the accompanying figures:

Figure 1 provides microscopy images of chondrocyte cell growth on gelatin beads (A) and PLGA beads (B) (Examples 8 and 10).

5 Figure 2 shows results of evaluation of cells for phenotype using RT-PCR, wherein PCR products are analysed by electrophoresis on 2% agarose gels (Example 20).

Figure 3 shows the effect of beads on gel contraction after a 2-week culture of chondrocytes with and without beads (gelatin) in a collagen type I gel (Example 28).

10 Figure 4 shows an example of new tissue formation using cultured chondrocytes on demineralised bone particles with a collagen type I gel (Example 31).

Example 1: Chondrocyte isolation

Fresh cartilage tissue is collected in DMEM/10% FBS or autologous serum
15 containing 100µg/ml penicillin and streptomycin. After weighing, the tissue is placed in a sterile petri dish containing 3-4 ml of DMEM and dissected into 1 mm³ pieces using a sharp sterile scalpel. It is then digested with 10% w/v trypsin in PBS at 37°C for 1 hour. Approximately 2ml of 10 % w/v trypsin is used per gram of tissue. The residual tissue pieces are collected by centrifugation (1000 rpm, 5 mins) and washed
20 with PBS, then water (using approximately 5-10 ml per gram of tissue). A second digestion step is then performed overnight at 37°C using 2 ml of a mixture of bacterial collagenase and hyaluronidase per gram of tissue. The digestion mixture is prepared by adding 2 mg hyaluronidase (1520 units) and 200 µl of collagenase stock (taken from a 3000 unit/ml stock, stored at -70°C in a buffer of 50 mM tris, 10 mM CaCl₂, pH 7.0) to
25 2 ml of DMEM and filter sterilising. The digested tissue is passed through a 70µm Nylon cell strainer and the cells are washed and collected by centrifugation. Cell numbers and viability are assessed using a trypan blue count on a small known aliquot.

Example 2: Fibroblast isolation

30 Fresh skin, after hair removal and washing in 70% ethanol, is collected in DMEM/10% FBS or autologous serum containing 100µg/ml penicillin and streptomycin. The tissue is placed in a sterile petri dish containing 3-4 ml of DMEM and dissected into 1 mm³ pieces using a sharp sterile scalpel. The tissue pieces are left in culture in DMEM/10% FBS or autologous serum containing 100µg/ml penicillin and
35 streptomycin to allow migration of fibroblasts onto the tissue culture plastic. After cells are visible on the tissue culture plastic, the tissue is removed and the cells sub-

cultured. Cell numbers and viability are assessed using a trypan blue count on a small known aliquot.

Example 3: Osteoblast isolation

- 5 Fresh cortical bone is collected in DMEM/10% FBS or autologous serum containing 100µg/ml penicillin and streptomycin. The bone is placed in a sterile petri dish containing 3-4 ml of DMEM. The bone piece(s) are left in culture in DMEM/10% FBS or autologous serum containing 100µg/ml penicillin and streptomycin to allow migration of osteoblasts onto the tissue culture plastic. After cells are visible on the
- 10 tissue culture plastic, the bone is removed and the cells sub-cultured. Cell numbers and viability are assessed using a trypan blue count on a small known aliquot.

Example 4: Stem cell isolation

- Adult mesenchymal stem cells (MSC) are harvested from bone marrow
- 15 aspirates. The marrow is washed twice with sterile PBS then resuspended in DMEM/10% FBS or autologous serum containing 100µg/ml penicillin and streptomycin. Marrow cells are then layered onto a Percoll cushion (1.073g/ml density) and cells collected after centrifugation for 30 min. at 250g and transferred to tissue culture flasks. Various additives including dexamethasone, growth factors and
- 20 cytokines are used to select and propagate specific cell lineages.

Example 5: Cell culture in monolayers

- Cells, such as fibroblasts, chondrocytes, osteoblasts and other types isolated according to the protocols described above in Examples 1-4, are cultured on tissue
- 25 culture plastic in DMEM/10% FBS or autologous serum containing 100µg/ml penicillin and streptomycin, at 37°C in 5% carbon dioxide atmosphere. Medium additions or change is performed every 2 days. Cells are grown to confluency, then trypsinised and replated into flasks as monolayers or transferred to beads/particles.

Example 6: Cell culture on non-resorbable beads

- Beads or particles, for example Cytodex beads (Pharmacia Biotech), providing a surface area of 250-500 cm², are pre-washed with 50 ml of warmed media (DMEM/10% FBS or autologous serum containing 100µg/ml penicillin and streptomycin) at 37°C then placed inside a 125 ml spinner bottle. 1 x 10⁵ cells, either
- 35 freshly isolated cells, previously passaged cells or previously isolated and frozen cells, are added to the beads or particles. The bottle is then stirred in a 37°C incubator (with

- 5% CO₂), at 25 rpm intermittently for 2 minutes every 30 minutes for 3 hours, then intermittently for 2 minutes every 30 minutes for the next 3 hours, then continuously first at 45rpm for 15 minutes, then 50rpm for 15 minutes, 55rpm for 15 minutes, then to the final speed of 60 rpm. The cells are then grown at this speed until 90 % confluence is achieved, usually 5-8 days depending on the original inoculum. For collection of the cells on the beads or particles, either for release and further seeding or for preparation for delivery to a patient or further processing, the cells and beads are washed with warm, 37°C PBS and collected by centrifugation.

10 **Example 7: Preparation of gelatin beads**

- Gelatin microparticles are synthesized by using emulsion method. Briefly, gelatin is dissolved in 50 mM acetic acid to 20% (w/v). Two hundred milliliters olive oil is warmed up to 37°C. The warmed olive oil is stirred at 300 rpm. Forty millilitres gelatin solution kept at 37°C is then applied to olive oil through a 20-gauge needle.
- 15 This solution is also prepared containing 10% w/w native collagen. The emulsion is kept stirred for 90 minutes. The emulsion is then cooled down by stirring at 4°C for 30 minutes in order to harden the gelatin particles. Five hundred millilitres of 0.2% Triton X-100 in PBS is added to the emulsion and stirred at room temperature for 10 minutes. The mixture is then put in a separating funnel and settled for one hour. The liquid in
- 20 the lower portion is collected and after gelatin microparticles precipitate, the upper liquid decanted off carefully and the particles rinsed with water two times. Five hundred millilitres of 0.1% glutaraldehyde in PBS is added to the gelatin microparticles and stirred for one hour for cross-linking. The cross-linked gelatin beads are then rinsed with water several times and soaked in ethanol. The ethanol is decanted and the
- 25 gelatin microparticles dried under vacuum. Before seeding cells, the gelatin beads are rehydrated with PBS overnight and then with chondrocyte medium. The average size of gelatin microparticles is about 110 µm.

Example 8: Cell-culture on gelatin beads

- 30 Gelatin beads, providing a surface area of 250-500 cm², are pre-washed with 50 ml of warmed media (DMEM / 10% FBS or autologous serum containing 100µg/ml penicillin and streptomycin) at 37°C then placed inside a 125 ml spinner bottle. 1 x 10⁵ cells, either freshly isolated cells, previously passaged cells or previously isolated and frozen cells, are added to the beads or particles. The bottle is then stirred in a 37°C
- 35 incubator (with 5% CO₂), at 25 rpm intermittently for 2 minutes every 30 minutes for 3 hours, then 45rpm intermittently for 2 minutes every 30 minutes for the next 3 hours,

then continuously first at 45rpm for 15 minutes, then 50rpm for 15 minutes, 55rpm for 15 minutes, then to the final speed of 60 rpm. The cells are then grown at this speed until 90 % confluence is achieved, usually 5-8 days depending on the original inoculum. For collection of the cells on the beads or particles, either for release and
5 further seeding or for preparation for delivery to a patient or further processing, the cells and beads are washed with warm, 37°C PBS and collected by centrifugation. Figure 1A shows cell growth on gelatin beads 7 days after addition of chondrocytes to the gelatin beads.

10 **Example 9: Preparation of PLGA beads and particles**

Poly(lactide-co-glycolide) 85:15 w/w (PLGA) was dissolved in tetrahydrofuran and then emulsified into an aqueous solution containing 1% polyvinylalcohol by stirring. PLGA beads were collected by allowing them to settle, and were washed 5
15 times with water by decantation. Beads were then dried in a vacuum over night. Beads in the range of 30 μm to 300 μm were typically obtained, with an average size of 105 μm . Beads were fractionated into a narrower size range, 80 μm to 120 μm , by sieving. Alternatively, PLGA particles in the desired size range were obtained by crushing larger particles in a homogeniser, using a suspension of 1 g PLGA in 500 ml of water. Sieving provided particles of irregular shape in the desired size range, for example 50
20 μm to 250 μm . Surface modification of the PLGA beads and particles was carried out by adsorption of collagen I or collagen II from a solution containing 50 $\mu\text{g/ml}$ collagen in phosphate buffered saline at room temperature for 1 hour. Subsequent washing in phosphate buffered saline removed loosely bound collagen.

25 **Example 10: Cell culture on PLGA beads**

PLGA beads providing a surface area of 250-500 cm^2 , are pre-washed with 50 ml of warmed media (DMEM / 10% FBS or autologous serum containing 100 $\mu\text{g/ml}$ penicillin and streptomycin) at 37°C then placed inside a 125 ml spinner bottle. 1×10^5 cells, either freshly isolated cells, previously passaged cells or previously isolated and
30 frozen cells, are added to the beads or particles. The bottle is then stirred in a 37°C incubator (with 5% CO_2), at 25 rpm intermittently for 2 minutes every 30 minutes for 3 hours, then 45rpm intermittently for 2 minutes every 30 minutes for the next 3 hours, then continuously first at 45rpm for 15 minutes, then 50rpm for 15 minutes, 55rpm for 15 minutes, then to the final speed of 60 rpm. The cells are then grown at this speed
35 until 90 % confluence is achieved, usually 5-8 days depending on the original inoculum. For collection of the cells on the beads or particles, either for release and

further seeding or for preparation for delivery to a patient or further processing, the cells and beads are washed with warm, 37°C PBS and collected by centrifugation. Figure 1B shows chondrocyte culture on PLGA beads 14 days after chondrocytes were added to the PLGA beads. The chondrocytes have been stained with goat anti-type II collagen antibodies thereby indicating type II collagen synthesis.

Example 11: Preparation of bone particles

Fresh bone, free from adherent tissue and rinsed with phosphate buffered saline (PBS) is dried and then crushed and milled to provide particles which are separated by sieving, to give for example a fraction that passes through a 120 micron sieve, but is retained by an 80 micron sieve. These particles are degreased by washing in methanol, dichloromethane and acetone. Particles are then washed in 2 changes of PBS and then water and dried. Demineralised bone particles are prepared by agitation of bone particles in 0.5 M EDTA, pH 7.4, for 20 hr. After separation by gentle centrifugation, this process was repeated at least a further two times.

Example 12: Cell culture on bone particles

Culture of cells on bone particles was as in Example 10, except bone particles, both untreated and demineralised, are used instead of PLGA beads.

Example 13: Cell culture in a bioreactor

Beads or particles with cells attached, as described in Examples 6 or 8 or 10 or 12, are placed in a bioreactor, such as a High Aspect Ratio Vessel of a Synthecon™ Rotary Cell Culture System, where the vessel is filled with DMEM / 10% FBS or autologous serum containing 100µg/ml penicillin and streptomycin and air bubbles removed. Culture is continued in a humidified incubator with 5% carbon dioxide present, with the initial rotation speed at 15 rpm. The speed is then further adjusted, dependent on the nature and size of the bead or particle so that the beads or particles are not settling nor colliding with the edge of the vessel, but are forming a fluid orbit within the culture vessel. Medium change or addition is every 1 or 2 days.

Example 14: Removal and transfer of cells from a monolayer culture

Warm, 37°C, 0.3 % w/v trypsin in PBS is added directly to tissue culture flask, 5ml per 25 cm². After standing for up to 5 minutes, cells are dislodged from the plastic by gentle pipette action or by gentle mechanical action. Cells in the trypsin solution are collected by centrifugation at 1000 rpm for 5mins. The supernatant is then removed

and the cells gently resuspended in 5ml of media. Cells are counted using a trypan blue method.

Example 15: Removal of cells from polymer beads

- 5 Apply 6 ml of warm 0.3 % w/v trypsin directly to the collected and washed cells on beads and incubate at 37°C for 10 to 15 minutes without stirring. Apply 20ml of warm PBS to the mixture and gently pipette up and down to dislodge cells from beads or particles, which have a size greater than 70 μ m. Transfer cells and beads or particles through a 70 μ m filter into a 50ml tube. Collect the cells that pass through the filter by
10 centrifugation at 1000 rpm for 5mins. Remove the supernatant and gently resuspend the cells in 5ml of media. Cells are counted using a trypan blue method.

Example 16: Removal of cells from gelatin beads

- 15 Apply 6 ml of warm 0.3 % w/v trypsin directly to the collected and washed cells on beads and incubate at 37°C for 20 minutes. The gelatin beads were digested by the enzyme, releasing the cells into solution without the need for extensive mechanical agitation. Cells were collected by centrifugation at 1000 rpm for 5mins. Remove the supernatant and gently resuspend the cells in 5ml of media. Cells are counted using a trypan blue method.

20

Example 17: Transfer of cells onto resorbable beads for implant

- Cells, such as fibroblasts, chondrocytes, osteoblasts or other types, either freshly isolated, or previously passaged in monolayer culture or on non-resorbable beads or particles or on resorbable beads or particles, or previously isolated, cultured and frozen,
25 are suspended in warmed media (DMEM / 10% FBS or autologous serum containing 100 μ g/ml penicillin and streptomycin) at 37°C, and added to pre-washed beads or particles, as in Examples 7 or 9 or 11, and attachment is by a gradual increase in agitation, as in Examples 6 or 8 or 10 or 12.

30 **Example 18: Evaluation of cells by alcian blue staining**

- An advantage of culturing cells on beads or particles (Example 6, 8, 10, 12) is the control of phenotype. For articular cartilage, the phenotype is monitored using a variety of histochemical and immunohistochemical markers that can distinguish chondrocytes from de-differentiated fibrochondrocytes. Alcian blue, a general stain for
35 the glycosaminoglycans of articular cartilage, is prepared as a 2% filtered solution in 3% acetic acid at pH 2.5. After fixing in neutral buffered formaldehyde for 2-3min,

slides are incubated in 3% acetic acid for 3min. Alcian blue solution is applied for at least 20hr at 37°C, slides are rinsed with water and a 2 minute neutral red stain is applied. An ethanol rinse is used prior to mounting in HistoClear.

5 **Example 19: Evaluation of cells by immunohistological staining**

The phenotype of cultured cells is monitored by specific immunological markers. For articular chondrocytes antibodies against collagen type II is used to monitor the correct phenotype and an anti-collagen type I antibody is used to monitor the extent of change or de-differentiation. If cells are to be stained for matrix
10 production, for example by anti-collagen antibodies, fresh ascorbic acid must be added to cultures daily to a final concentration of 50µg/ml for at least 6 days. After washing in warm PBS, cells on beads are pre-fixed, once in 50 % (v/v) methanol in PBS for 10 minutes, twice in cool 70 % (v/v) methanol in PBS for 10 minutes, then finally in 70 % (v/v) ethanol in H₂O. Formalin or glutaraldehyde may be used as alternative fixatives
15 for use with proteoglycans stains such as Alcian Blue. The primary antibody is diluted in PBS (e.g. goat anti type II collagen diluted 1 in 5 with PBS) and is applied for 1 hr at room temperature, then, after washing with PBS, an FITC-conjugated antibody diluted in PBS (e.g. rabbit anti goat FITC diluted 1 in 200 with PBS) is applied for 1 hr at room temperature. After washing with PBS twice, the beads are resuspended in mounting
20 medium (e.g. 90% glycerol, 10% PBS, 0.025 % DABCO). Fluorescent images are collected on an Optiscan confocal microscope.

Example 20: Evaluation of cells by in situ hybridisation and RT-PCR

Cells for *in situ* hybridisation characterisation are fixed as in Example 19. *In situ*-hybridization for mRNA encoding, for example collagen type I or collagen type II
25 is performed using UTP-³³P detection following the method of Bisucci T, Hewitson TD, Darby IA, (2000) "cRNA probes: comparison of isotopic and non-isotopic detection methods", in Methods in Molecular Biology, 123: 291-303. A type I collagen riboprobe consisting of 372 bp region of the human collagen pro $\alpha 1$ (I) gene or a type II
30 collagen riboprobe consisting of a 200 bp region of the bovine collagen $\alpha 1$ (II) gene, is used.

For RT-PCR cells (pig chondrocytes) are cultured in monolayers and retrieved as in Example 5 and Example 14. Cells are lysed thoroughly in 1 ml REzol™ C&T (USA) by vortexing. The cell lysate is transferred to a microfuge tube, and incubated
35 for 5 minutes at room temperature. Cell lysate is then mixed vigorously with 0.2 ml of chloroform and incubated at room temperature for 2 minutes. After centrifugation at

12,000 x g for 15 minutes at 4°C, the upper aqueous layer is transferred to a new microfuge, and an equal volume of isopropanol is added and mixed gently. The samples were incubated at room temperature for 10 minutes and centrifuged at 12,000 x g for 10 minutes at 4°C. The supernatant is removed carefully, and the RNA pellet is washed in 1 ml of 75% ethanol by vortex mixing and then centrifuged at 12,000 x g for 5 minutes at 4°C. The ethanol is then removed carefully and the RNA pellet dried by air. The RNA pellet is dissolved in 20 µl of DEPC-treated water. The mRNA is then reverse-transcribed into cDNA by using oligo-dT primer and SUPERSRIPT™II following manufacturer's recommendations (Life Technologies).

10 Aliquots of 2 µl from the RT reactions are used for amplification of transcripts using primers specific for the analyzed genes. PCR reactions are carried out by 3 minutes denaturation at 95°C, followed by 35 cycles of 1 minute denaturation at 95°C, 1 minute annealing at 50°C and 1 minute elongation at 72°C. The primers for analyzed genes are designed as following:

15

β-actin:	5' -AACGGCTCCGGCATGTGC-3' (SEQ ID NO:1) and
	5' -GGGCAGGGGTGTTGAAGG-3' (SEQ ID NO:2)
20 Type I collagen:	5' -GCTGGCCAACTATGCCTC-3' (SEQ ID NO:3) and
	5' -GAAACAGACTGGGCCAATG-3' (SEQ ID NO:4)
Type II collagen:	5' -TGCTTACCTGGACGAAGC-3' (SEQ ID NO:5) and
25	5' -CCCAGTTCAGGCTCTTAG-3' (SEQ ID NO:6)
SOX9:	5' -CCCAACGCCATCTTCAAG-3' (SEQ ID NO:7) and
	5' -CTTGGACATCCACACGTG-3' (SEQ ID NO:8)
30 Aggrecan:	5' -CTGTTACCGCCACTTCCC-3' (SEQ ID NO:9) and
	5' -GGTGCGGTACCAGTGCAC-3' (SEQ ID NO:10)

This is shown in Figure 2.

Example 21: Synthetic gel preparation

A suitable gel, that is bioresorbable, is formed by using a precursor consisting of PEO polymerised at its termini with oligomers of α -hydroxy acids, such as glycolic acid or lactic acid, and end capped at all oligo(α -hydroxy acid) termini with a polymerisable acrylate group, allowing polymerisation of the precursor to form a gel by brief exposure to long wavelength ultraviolet light.

Example 22: Preparation of a cells and beads and synthetic gel mixture

Cells, after removal from a gelatin bead substrate as shown in Example 8, or from other substrates, are mixed with fresh gelatin beads, made as in Example 7, or other bioresorbable beads or particles as in Example 9 or Example 11, in DMEM containing autologous serum or bovine fetal calf serum, and mixed with a synthetic gel precursor, such as that of Example 21, to form a uniform mixture, with the gel being formed by a brief exposure to ultraviolet light.

Example 23: Biological (collagen) gel preparation

Four grams of type I collagen, type II collagen, or mixtures of these collagens were dissolved in 1 litre 50 mM acetic acid solution. The collagen solution was spun at 9500 rpm, 4°C for 45 minutes. The supernatant was collected. The collagen solution was put into a dialysis bag and then dialyzed against 25 litres 1M acetic acid for two days, then against 25 litres water for four days with multiple water changes. The collagen solution was then concentrated in the sealed dialysis bag by hanging in a laminar flow hood for a day. The final concentration of the collagen solution was about 20 mg/ml (2% w/v).

Example 24: Preparation of a cells, beads and biological gel mixture

Cells, after removal from a gelatin bead substrate as shown in Example 8, or from other substrates, are mixed with fresh gelatin beads, made as in Example 7, or other bioresorbable beads or particles, in DMEM containing autologous serum or bovine fetal calf serum, and mixed with a biological gel or precursor, such as a 2% collagen solution prepared as in Example 23, to form a uniform mixture with the cells and beads or particles uniformly mixed, with gel formation being achieved by incubation of the mixture at 37°C.

Example 25: Preparation of cells-on-beads and a synthetic gel mixture

Cells attached to a gelatin bead substrate as shown in Example 8, or to other bioresorbable beads or particles, are collected by allowing the culture mixture to settle, with the excess culture media then being removed. The cells on the beads are then mixed with a synthetic gel precursor, such as that of Example 21, to form a uniform mixture, with the gel being formed by a brief exposure to ultraviolet light.

Example 26: Preparation of cells-on-beads and a biological gel mixture

Cells attached to a gelatin bead substrate as shown in Example 8, or to other bioresorbable beads or particles, are collected by allowing the culture mixture to settle, with the excess culture media then being removed. The cells on the beads are then mixed with a biological gel or precursor, such as a 2% collagen solution prepared as in Example 23, to form a uniform mixture. Nine parts of the collagen solution was mixed with one part of 10 X DMEM and 0.1 part of 1N NaOH. Four parts of this mixture was mixed 1 part of chondrocyte-gelatin bead composites. Gel formation was achieved by incubation at 37°C incubator for an hour, or could be achieved by body temperature for an implanted mixture.

Example 27: In vitro culture of a cells/beads/biological gel mixture

A biological gel containing cells and beads, as prepared in Example 24, is transferred, for example to a 24-well plate, and 1.5 ml of chondrocyte medium is added to each sample. Chondrocyte medium is changed every other day and 100 µg/ml of ascorbic acid is supplied every day. For *in vitro* evaluation, samples are collected after 3 days, 7 days, 14 days, 21 days and 28 days.

Example 28: In vitro culture of a cell-on-beads/biological gel mixture

A biological gel containing cells-on-beads, as prepared in Example 26, is transferred to a cell culture plate and cultured in the presence of ascorbic acid as described in Example 27. Chondrocytes associated with the beads proliferate in the gel by day 3 and secreted new matrix of collagen type II and glycosaminoglycans consistent with the chondrocyte phenotype. The presence of the beads substantially reduces the rate and extent of gel contraction as shown in Figure 3.

Example 29: In vitro culture of a cells/beads/synthetic gel mixture

A synthetic gel containing cells and beads, as prepared in Example 22, is transferred to a cell culture plate and cultured in the presence of ascorbic acid as described in Example 27.

Example 30: In vitro culture of a cells-on-beads/synthetic gel mixture

A synthetic gel containing cells on beads, as prepared in Example 25, is transferred to a cell culture plate and cultured in the presence of ascorbic acid as described in Example 27.

Example 31: Implant of a cells/beads/biological gel mixture into animals

Either a cells and beads or a cells-on-beads in a type I collagen gel, as shown in Example 24 or 26, is injected subcutaneously into nude mice. Sacrifice of animals after 1 month and 2 months allows histological and immunohistological evaluation of the new tissue formed. Explants from nude mice show that articular cartilage can be produced using a variety of beads including gelatin, modified gelatin with collagen type I, and demineralised bone. Using type I collagen as the delivery gel, good tissue formation is noted within 1 month and continued at 2 months. Histochemical and immunohistochemical evaluation as described in Examples 18, 19 and 20 demonstrates the correct matrix and cartilage phenotype. Figure 4 shows an example of new tissue formation using cultured chondrocytes on demineralised bone particles with a collagen type I gel.

Example 32: Implant of an in vitro cultured material into animals

Either a cells and beads or a cells-on-beads in a biological gel mixture, for example using fibroblasts, chondrocytes or osteoblasts and gelatin beads in a type I collagen gel, as shown in Example 27 or 28 is surgically implanted subcutaneously into nude mice. Sacrifice of animals after 1 month and 2 months allows histological evaluation of the new tissue formed.

Example 33: Implant of a cells-on-beads/synthetic gel mixture into animals

Either a cells and beads or a cells-on-beads in a synthetic gel mixture, for example a polyethylene glycol/lactic-glycolic acid/ α -hydroxy acid type as shown in Example 22 or 25 is injected subcutaneously into nude mice. Sacrifice of animals after 1 month and 2 months allows histological evaluation of the new tissue formed.

Example 34: Repair of a cartilage defect using a cell containing mixture

A preparation of cells (chondrocytes) and beads or particles and a gel is used. This mixture, for example chondrocytes attached to a gelatin bead substrate in a 2% type I collagen mixture, as shown in Example 26, is loaded into a syringe with a needle of sufficient diameter to allow easy passage of the beads or particles, such as 22 gauge. The material is then injected into a cartilage defect established in the knee of a sheep. The implanted material may also be retained in place by affixing a piece of autologous periosteum over the implanted chondrocyte containing material. After closure of the wound, the knee is kept temporally immobile to allow the collagen to form a semi-solid gel.

Example 35: Repair of a cartilage defect using a cell containing mixture

Repair of a knee defect using a preparation of cells (chondrocytes) and beads or particles and a gel is achieved as shown in Example 34, except that a synthetic gel, as shown in Example 21 is used, with gel formation being achieved once the material is in the cartilage defect by brief exposure to ultraviolet light. The implanted material may also be retained in place by affixing a piece of autologous periosteum over the implanted chondrocyte containing material.

Example 36: Repair of a cartilage defect using an in vitro cultured implant

A preparation of cells (chondrocytes) and beads or particles and a gel is used. This mixture, for example chondrocytes attached to a gelatin bead substrate in a 2% type I collagen mixture, as shown in Example 27, is held in cell culture supplemented by ascorbic acid for 10 days to allow a tissue like material to form containing the chondrocytes and gelatin beads. The tissue like material is then surgically implanted into a cartilage defect established in the knee of a sheep. The implanted material may also be retained in place by affixing a piece of autologous periosteum over the implanted chondrocyte containing material.

Example 37: Repair of a bone defect using a cell containing mixture

A material is prepared as in Example 34, but with osteoblasts as the cell component and crushed bone particles, and is injected into a round defect in a sheep femur. Histological examination after 2 months is used to demonstrate bone repair.

Example 38: Repair of a bone defect using a cell containing mixture

A material containing osteoblasts, crushed bone particles and type I collagen is prepared as in Example 37, but with the addition of BMP 2 or other growth factors.

- 5 The material is injected into a round defect in a sheep femur and examined by histology after 2 months to demonstrate bone repair.

Example 39: Repair of a tissue defect using a cell containing mixture

- 10 A material is prepared as in Example 34, but with fibroblasts as the cell component and gelatin beads, and is injected subcutaneously into sheep. Histological examination after 2 months is used to demonstrate tissue repair.

Example 40: Repair of a tissue defect using a cell containing mixture

- 15 A material is prepared as in Example 34, but with adipocytes as the cell component and gelatin beads, and is injected subcutaneously into sheep. Histological examination after 2 months is used to demonstrate tissue repair.

Example 41: Repair of a tissue defect using a cell containing mixture

- 20 A material is prepared with two cell types, fibroblasts and adipocytes, as the cell component, cultured separately on gelatin beads, as in Examples 39 and 40, which are mixed in the collagen gel, and injected subcutaneously into sheep. Histological examination after 2 months is used to demonstrate tissue repair.

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It will be appreciated by persons skilled in the art that numerous variations and/or modifications may be made to the invention as shown in the specific embodiments without departing from the spirit or scope of the invention as broadly described. The present embodiments are, therefore, to be considered in all respects as
5 illustrative and not restrictive.

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Claims:

1. A method for treating diseased or damaged tissue in a subject, said method comprising administering to said subject at a site wherein said diseased or damaged
5 tissue occurs, cells of a type(s) normally found in healthy tissue corresponding to said diseased or damaged tissue, and/or suitable progenitor cells thereof, in association with bioresorbable beads or particles and optionally a gel and/or gel-forming substance.
2. The method of claim 1, wherein the cells and/or progenitor cells are associated
10 with the beads or particles by being bound thereto.
3. A method of claim 1 or 2, wherein the bioresorbable beads or particles are comprised of a pharmaceutically acceptable polymer(s).
- 15 4. The method of claim 3, wherein said polymer(s) is/are a biologically-based polymer(s) selected from the group consisting of gelatin and collagen.
5. The method of claim 3, wherein the polymer(s) is/are a synthetic polymer(s) selected from the group consisting of poly(glycolide), poly(lactide) and poly(lactide-
20 co-glycolide).
6. The method of claim 3, wherein said polymer(s) is a mixture of a biologically-based polymer(s) and a synthetic polymer(s), wherein said biologically-based polymer(s) is/are selected from the group consisting of gelatin and collagen, and said
25 synthetic polymer(s) is/are selected from the group consisting of poly(glycolide), poly(lactide) and poly(lactide-co-glycolide).
7. The method of claim 1 or 2, wherein the bioresorbable beads or particles are comprised of a pharmaceutically acceptable non-polymeric substance(s).
30
8. The method of claim 7, wherein the non-polymeric substance(s) is/are selected from the group consisting of crushed bone and demineralised bone.
9. The method of any one of claims 3 to 8, wherein said bioresorbable beads or
35 particles have been functionalised or coated in a suitable cell adherence-enhancing material.

10. The method of any one of claims 3 to 9, wherein said bioresorbable beads or particles are further comprised of a beneficial agent(s) selected from the group consisting of growth factors, glycosaminoglycans and hydrophilic compounds.
- 5 11. The method of any one of claims 1 to 10, wherein said bioresorbable beads or particles have a diameter or dimension sized in the range of about 20 to 2500 μm .
12. The method of claim 11, wherein the average size of said bioresorbable beads or
10 particles is about 50 to 200 μm .
13. The method of any one of claims 1 to 12, wherein said gel and/or gel-forming substance is bioresorbable.
- 15 14. The method of claim 13, wherein said gel and/or gel-forming substance comprises a biologically-based polymer(s) selected from the group consisting of collagen, fibrin, hyaluronan, chitosan and mixtures thereof.
- 15 15. The method of claim 13, wherein said gel and/or gel-forming substance
20 comprises a synthetic polymer(s) selected from the group consisting of photopolymerizable end-capped block copolymers of poly(ethylene oxide) and an α -hydroxy acid.
16. The method of claim 13, wherein said gel and/or gel-forming substance
25 comprises a mixture of a biologically-based polymer(s) and a synthetic polymer(s), wherein said biologically-based polymer(s) is/are selected from the group consisting of collagen, fibrin, hyaluronan, chitosan and mixtures thereof, and said synthetic polymer(s) is/are selected from the group consisting of photopolymerizable end-capped block copolymers of poly(ethylene oxide) and an α -hydroxy acid.
- 30 17. The method of any one of claims 13 to 16, wherein said gel and/or gel-forming substance is further comprised of a beneficial agent(s) selected from the group consisting of growth factors, glycosaminoglycans and hydrophilic compounds.
- 35 18. The method of any one of claims 13 to 17, wherein said gel and/or gel-forming substance includes an adhesive material(s).

19. The method of any one of claims 1 to 18, wherein an average of between about 3 and 500 cells and/or progenitor cells are associated with each of said beads or particles.
- 5 20. The method of any one of claims 1 to 19, wherein said cells and/or progenitor cells are chondrocytes, embryonic stem cells and/or bone marrow stromal cells.
21. The method of any one of claims 1 to 19, wherein said cells and/or progenitor cells are fibroblast and/or progenitor cells thereof.
- 10 22. The method of any one of claims 1 to 19, wherein said cells and/or progenitor cells are adipocytes and/or progenitor cells thereof.
23. The method of any one of claims 1 to 19, wherein said cells and/or progenitor cells are osteoblasts and/or progenitor cells thereof.
- 15 24. The method of any one of claims 1 to 19, wherein said cells and/or progenitor cells are a mixture of cell types and/or progenitor cell types.
- 20 25. The method of any one of claims 1 to 19, wherein said cells and/or suitable progenitor cells thereof in association with said bioresorbable beads or particles and a gel and/or gel-forming substance, is administered by entrapping the gel and/or gel-forming substance within or under a tissue at said site where diseased or damaged tissue occurs.
- 25 26. The method of any one of claims 1 to 19, wherein said cells and/or suitable progenitor cells thereof in association with said bioresorbable beads or particles and a gel and/or gel-forming substance, is administered by entrapping the gel and/or gel-forming substance under a tissue flap or other membranous flap at said site where
- 30 diseased or damaged tissue occurs.
27. The method of claim 25 or 26, wherein said cells and/or suitable progenitor cells are chondrocytes and the diseased or damaged tissue to be treated is articular cartilage.
- 35 28. A method for treating disease or damaged tissue in a subject, said method comprising the steps of;

- (i) obtaining cells of a type(s) normally found in healthy tissue corresponding to said diseased or damaged tissue and/or suitable progenitor cells thereof,
 - (ii) expanding said cells and/or progenitor cells in the presence of bioresorbable beads or particles whereby said expanded cells and/or progenitor cells
5 become bound to the said beads or particles, and
 - (iii) administering to said subject the beads or particles with said cells and/or progenitor cells bound thereto optionally in a gel and/or gel-forming substance at a site wherein said diseased or damaged tissue occurs.
- 10 29. The method of claims 28, wherein step (ii) is conducted in a bioreactor containing a suitable culture medium, and wherein said culture medium is agitated and aerated.
- 15 30. The method of claim 29, wherein said bioreactor is a tumbler-type bioreactor equipped with internal veins to assist in movement of the cells and/or progenitor cells, culture medium and bioresorbable beads or particles.
31. The method of claim 29, wherein said bioreactor is a spinner flask.
- 20 32. A method for the treatment of diseased or damage tissue in a subject, said method comprising the steps of,
- (i) obtaining cells of a type(s) normally found in healthy tissue corresponding to said diseased or damaged tissue and/or suitable progenitor cells thereof,
 - (ii) expanding said cells and/or progenitor cells,
 - 25 (iii) binding said expanded cells and/or progenitor cells to bioresorbable beads or particles, and
 - (iv) administering to said subject the beads or particles with said cells and/or progenitor cells bound thereto optionally in a gel and/or gel-forming substance at a site wherein said diseased or damaged tissue occurs.
- 30 33. The method of any one of claims 28 to 32, wherein the bioresorbable beads or particles are comprised of a pharmaceutically acceptable polymer(s).
34. The method of claim 33, wherein said polymer(s) is/are a biologically-based
35 polymer(s) selected from the group consisting of gelatin and collagen.

35. The method of claim 33, wherein the polymer(s) is/are a synthetic polymer(s) selected from the group consisting of poly(glycolide), poly(lactide) and poly(lactide-co-glycolide).

5 36. The method of claim 33, wherein said polymer(s) is a mixture of a biologically-based polymer(s) and a synthetic polymer(s), wherein said biologically-based polymer(s) is/are selected from the group consisting of gelatin and collagen, and said synthetic polymer(s) is/are selected from the group consisting of poly(glycolide), poly(lactide) and poly(lactide-co-glycolide).

10

37. The method of any one of claims 28 to 32, wherein the bioresorbable beads or particles are comprised of a pharmaceutically acceptable non-polymeric substance(s).

38. The method of claim 37, wherein the non-polymeric substance(s) is/are selected
15 from the group consisting of crushed bone and demineralised bone.

39. The method of any one of claims 28 to 38, wherein said bioresorbable beads or particles have been functionalised or coated in a suitable cell adherence-enhancing material.

20

40. The method of any one of claims 28 to 39, wherein said bioresorbable beads or particles are further comprised of a beneficial agent(s) selected from the group consisting of growth factors, glycosaminoglycans and hydrophilic compounds.

25 41. The method of any one of claims 28 to 40, wherein said bioresorbable beads or particles have a diameter or dimension sized in the range of about 20 to 2500 μm .

42. The method of claim 41, wherein the average size of said bioresorbable beads or particles is about 50 to 200 μm .

30

43. The method of any one of claims 28 to 42, wherein said gel and/or gel-forming substance is bioresorbable.

44. The method of claim 43, wherein said gel and/or gel-forming substance
35 comprises a biologically-based polymer(s) selected from the group consisting of collagen, fibrin, hyaluronan, chitosan and mixtures thereof.

45. The method of claim 43, wherein said gel and/or gel-forming substance comprises a synthetic polymer(s) selected from the group consisting of photopolymerizable end-capped block copolymers of poly(ethylene oxide) and an α -hydroxy acid.

46. The method of claim 43, wherein said gel and/or gel-forming substance comprises a mixture of a biologically-based polymer(s) and a synthetic polymer(s), wherein said biologically-based polymer(s) is/are selected from the group consisting of collagen, fibrin, hyaluronan, chitosan and mixtures thereof, and said synthetic polymer(s) is/are selected from the group consisting of photopolymerizable end-capped block copolymers of poly(ethylene oxide) and an α -hydroxy acid.

47. The method of any one of claims 43 to 46, wherein said gel and/or gel-forming substance is further comprised of a beneficial agent(s) selected from the group consisting of growth factors, glycosaminoglycans and hydrophilic compounds.

48. The method of any one of claims 43 to 47, wherein the gel and/or gel-forming substance includes an adhesive material(s).

49. The method of any one of claims 28 to 48, wherein an average of between about 3 and 500 cells and/or progenitor cells are bound to each of said beads or particles.

50. The method of any one of claims 28 to 49, wherein said cells and/or progenitor cells are chondrocytes, embryonic stem cells and/or bone marrow stromal cells.

51. The method of any one of claims 28 to 49, wherein said cells and/or progenitor cells are fibroblast and/or progenitor cells thereof.

52. The method of any one of claims 28 to 49, wherein said cells and/or progenitor cells are adipocytes and/or progenitor cells thereof.

53. The method of any one of claims 28 to 49, wherein said cells and/or progenitor cells are osteoblasts and/or progenitor cells thereof.

54. The method of any one of claims 28 to 49, wherein said cells and/or progenitor cells are a mixture of cell types and/or progenitor cell types.

55. The method of any one of claims 28 to 54, wherein step (ii) expands the cells
5 and/or progenitor cells 5 to 2000-fold.

56. The method of claim 55, wherein step (ii) expands the cells and/or progenitor cells 10 to 100-fold.

10 57. The method of any one of claims 28 to 49, wherein said cells and/or suitable progenitor cells thereof bound to said bioresorbable beads or particles and a gel and/or gel-forming substance, is administered by entrapping the gel and/or gel-forming substance within or under a tissue at said site where diseased or damaged tissue occurs.

15 58. The method of any one of claims 28 to 49, wherein said cells and/or suitable progenitor cells thereof in association with said bioresorbable beads or particles and a gel and/or gel-forming substance, is administered by entrapping the gel and/or gel-forming substance under a tissue flap or other membranous flap at said site where diseased or damaged tissue occurs.

20

59. The method of claim 57 or 58, wherein said cells and/or suitable progenitor cells are chondrocytes and the diseased or damaged tissue to be treated is articular cartilage.

60. The method of any one of the preceding claims, wherein the subject is a human
25 subject.

61. A device having tissue-like characteristics for treating diseased or damaged tissue in a subject, wherein said device comprises cells of a type(s) normally found in healthy tissue corresponding to said diseased or damaged tissue, and/or suitable
30 progenitor cells thereof, in association with bioresorbable beads or particles and optionally a gel and/or gel-forming substance.

62. A device having tissue-like characteristics for augmenting tissue in a subject, wherein said device comprises cells of a type(s) normally found in the tissue to be
35 augmented, and/or suitable progenitor cells thereof, in association with bioresorbable beads or particles and optionally a gel and/or gel-forming substance.

63. The device of claim 61 or 62, wherein the cells and/or progenitor cells are associated with the beads or particles by being bound thereto.
- 5 64. The device of any one of claims 61 to 63, wherein the bioresorbable beads or particles are comprised of a pharmaceutically acceptable polymer(s).
65. The device of claim 64, wherein said polymer(s) is/are a biologically-based polymer(s) selected from the group consisting of gelatin and collagen.
- 10 66. The device of claim 64, wherein the polymer(s) is/are a synthetic polymer(s) selected from the group consisting of poly(glycolide), poly(lactide) and poly(lactide-co-glycolide).
- 15 67. The device of claim 64, wherein said polymer(s) is a mixture of a biologically-based polymer(s) and a synthetic polymer(s), wherein said biologically-based polymer(s) is/are selected from the group consisting of gelatin and collagen, and said synthetic polymer(s) is/are selected from the group consisting of poly(glycolide), poly(lactide) and poly(lactide-co-glycolide).
- 20 68. The device of any one of claims 61 to 63, wherein the bioresorbable beads or particles are comprised of a pharmaceutically acceptable non-polymeric substance(s).
69. The device of claim 68, wherein the non-polymeric substance(s) is/are selected
25 from the group consisting of crushed bone and demineralised bone.
70. The device of any one of claims 64 to 69, wherein said bioresorbable beads or particles have been functionalised or coated in a suitable cell adherence-enhancing material.
- 30 71. The device of any one of claims 61 to 70, wherein said bioresorbable beads or particles are further comprised of a beneficial agent(s) selected from the group consisting of growth factors, glycosaminoglycans and hydrophilic compounds.
- 35 72. The device of any one of claims 61 to 70, wherein said bioresorbable beads or particles have a diameter or dimension sized in the range of about 20 to 2500 μm .

73. The device of claim 72, wherein the average size of said bioresorbable beads or particles is about 50 to 200 μm .

5 74. The device of any one of claims 61 to 73, wherein said gel and/or gel-forming substance is bioresorbable.

75. The device of claim 74, wherein said gel and/or gel-forming substance comprises a biologically-based polymer(s) selected from the group consisting of
10 collagen, fibrin, hyaluronan, chitosan and mixtures thereof.

76. The device of claim 75, wherein said gel and/or gel-forming substance comprises a synthetic polymer(s) selected from the group consisting of photopolymerizable end-capped block copolymers of poly(ethylene oxide) and an α -
15 hydroxy acid.

77. The device of claim 74, wherein said gel and/or gel-forming substance comprises a mixture of a biologically-based polymer(s) and a synthetic polymer(s), wherein said biologically-based polymer(s) is/are selected from the group consisting of
20 collagen, fibrin, hyaluronan, chitosan and mixtures thereof, and said synthetic polymer(s) is/are selected from the group consisting of photopolymerizable end-capped block copolymers of poly(ethylene oxide) and an α -hydroxy acid.

78. The device of any one of claims 61 to 77, wherein said gel and/or gel-forming
25 substance is further comprised of a beneficial agent(s) selected from the group consisting of growth factors, glycosaminoglycans and hydrophilic compounds.

79. The device of any one of claims 61 to 78, wherein said gel and/or gel-forming substance includes an adhesive material(s).
30

80. The device of any one of claims 61 to 79, wherein an average of between about 3 and 500 cells and/or progenitor cells are associated with each of said beads or particles.

35 81. The device of any one of claims 61 to 80, wherein said cells and/or progenitor cells are chondrocytes, embryonic stem cells and/or bone marrow stromal cells.

82. The device of any one of claims 61 to 80, wherein said cells and/or progenitor cells are fibroblast and/or progenitor cells thereof.
- 5 83. The device of any one of claims 61 to 80, wherein said cells and/or progenitor cells are adipocytes and/or progenitor cells thereof.
84. The device of any one of claims 61 to 80, wherein said cells and/or progenitor cells are osteoblasts and/or progenitor cells thereof.
- 10 85. The device of any one of claims 61 to 80, wherein said cells and/or progenitor cells are a mixture of cell types and/or progenitor cell types.
86. A method for treating diseased or damaged tissue in a subject, said method
15 comprising implanting into said subject at a site wherein said diseased or damaged tissue occurs a device having tissue-like characteristics, wherein said device comprises cells of a type(s) normally found in healthy tissue corresponding to said diseased or damaged tissue, and/or suitable progenitor cells thereof, in association with bioresorbable beads or particles and optionally a gel and/or gel-forming substance..
- 20 87. A method for augmenting tissue in a subject, said method comprising implanting into said subject at a site where tissue is to be augmented, a device having tissue-like characteristics, wherein said device comprises cells of a type(s) normally found in the tissue to be augmented, and/or suitable progenitor cells thereof, in
25 association with bioresorbable beads or particles and optionally a gel and/or gel-forming substance.
88. The method of claim 86 or 87, wherein the cells and/or progenitor cells are associated with the beads or particles by being bound thereto.
- 30 89. The method of any one of claims 86 to 88, wherein the bioresorbable beads or particles are comprised of a pharmaceutically acceptable polymer(s).
90. The method of claim 89, wherein said polymer(s) is/are a biologically-based
35 polymer(s) selected from the group consisting of gelatin and collagen.

91. The method of claim 89, wherein the polymer(s) is/are a synthetic polymer(s) selected from the group consisting of poly(glycolide), poly(lactide) and poly(lactide-co-glycolide).

5 92. The method of claim 89, wherein said polymer(s) is a mixture of a biologically-based polymer(s) and a synthetic polymer(s), wherein said biologically-based polymer(s) is/are selected from the group consisting of gelatin and collagen, and said synthetic polymer(s) is/are selected from the group consisting of poly(glycolide), poly(lactide) and poly(lactide-co-glycolide).

10

93. The method of any one of claims 86 to 88, wherein the bioresorbable beads or particles are comprised of a pharmaceutically acceptable non-polymeric substance(s).

94. The method of claim 93, wherein the non-polymeric substance(s) is/are selected
15 from the group consisting of crushed bone and demineralised bone.

95. The method of any one of claims 89 to 94, wherein said bioresorbable beads or particles have been functionalised or coated in a suitable cell adherence-enhancing material.

20

96. The method of any one of claims 86 to 95, wherein said bioresorbable beads or particles are further comprised of a beneficial agent(s) selected from the group consisting of growth factors, glycosaminoglycans and hydrophilic compounds.

25 97. The method of any one of claims 86 to 95, wherein said bioresorbable beads or particles have a diameter or dimension sized in the range of about 20 to 2500 μm .

98. The method of claim 97, wherein the average size of said bioresorbable beads or particles is about 50 to 200 μm .

30

99. The method of any one of claims 86 to 98, wherein said gel and/or gel-forming substance is bioresorbable.

100. The method of claim 99, wherein said gel and/or gel-forming substance
35 comprises a biologically-based polymer(s) selected from the group consisting of collagen, fibrin, hyaluronan, chitosan and mixtures thereof.

101. The method of claim 100, wherein said gel and/or gel-forming substance comprises a synthetic polymer(s) selected from the group consisting of photopolymerizable end-capped block copolymers of poly(ethylene oxide) and an α -
5 hydroxy acid.

102. The method of claim 99, wherein said gel and/or gel-forming substance comprises a mixture of a biologically-based polymer(s) and a synthetic polymer(s), wherein said biologically-based polymer(s) is/are selected from the group consisting of
10 collagen, fibrin, hyaluronan, chitosan and mixtures thereof, and said synthetic polymer(s) is/are selected from the group consisting of photopolymerizable end-capped block copolymers of poly(ethylene oxide) and an α -hydroxy acid.

103. The method of any one of claims 86 to 102, wherein said gel and/or gel-forming
15 substance is further comprised of a beneficial agent(s) selected from the group consisting of growth factors, glycosaminoglycans and hydrophilic compounds.

104. The method of any one of claims 86 to 103, wherein said gel and/or gel-forming substance includes an adhesive material(s).
20

105. The method of any one of claims 86 to 104, wherein an average of between about 3 and 500 cells and/or progenitor cells are associated with each of said beads or particles.

25 106. The method of any one of claims 86 to 105, wherein said cells and/or progenitor cells are chondrocytes, embryonic stem cells and/or bone marrow stromal cells.

107. The method of any one of claims 86 to 105, wherein said cells and/or progenitor cells are fibroblast and/or progenitor cells thereof.
30

108. The method of any one of claims 86 to 105, wherein said cells and/or progenitor cells are adipocytes and/or progenitor cells thereof.

109. The method of any one of claims 86 to 105, wherein said cells and/or progenitor
35 cells are osteoblasts and/or progenitor cells thereof.

110. The method of any one of claims 86 to 105, wherein said cells and/or progenitor cells are a mixture of cell types and/or progenitor cell types.

111. The method of any one of claims 86 to 110, wherein said subject is a human
5 subject.

112. A method for augmenting tissue in a subject, said method comprising administering to said subject at a site where tissue is to be augmented, cells of a type(s) normally found in the tissue to be augmented, and/or suitable progenitor cells thereof,
10 in association with bioresorbable beads or particles and optionally a gel and/or gel-forming substance.

113. The method of claim 112, wherein the cells and/or progenitor cells are associated with the beads or particles by being bound thereto.
15

114. A method of claim 112 or 113, wherein the bioresorbable beads or particles are comprised of a pharmaceutically acceptable polymer(s).

115. The method of claim 114, wherein said polymer(s) is/are a biologically-based
20 polymer(s) selected from the group consisting of gelatin and collagen.

116. The method of claim 114, wherein the polymer(s) is/are a synthetic polymer(s) selected from the group consisting of poly(glycolide), poly(lactide) and poly(lactide-co-glycolide).
25

117. The method of claim 114, wherein said polymer(s) is a mixture of a biologically-based polymer(s) and a synthetic polymer(s), wherein said biologically-based polymer(s) is/are selected from the group consisting of gelatin and collagen, and said synthetic polymer(s) is/are selected from the group consisting of poly(glycolide),
30 poly(lactide) and poly(lactide-co-glycolide).

118. The method of claim 112 or 113, wherein the bioresorbable beads or particles are comprised of a pharmaceutically acceptable non-polymeric substance(s).

35 119. The method of claim 118, wherein the non-polymeric substance(s) is/are selected from the group consisting of crushed bone and demineralised bone.

120. The method of any one of claims 114 to 119, wherein said bioresorbable beads or particles have been functionalised or coated in a suitable cell adherence-enhancing material.

5

121. The method of any one of claims 114 to 120, wherein said bioresorbable beads or particles are further comprised of a beneficial agent(s) selected from the group consisting of growth factors, glycosaminoglycans and hydrophilic compounds.

10 122. The method of any one of claims 112 to 121, wherein said bioresorbable beads or particles have a diameter or dimension sized in the range of about 20 to 2500 μm .

123. The method of claim 122, wherein the average size of said bioresorbable beads or particles is about 50 to 200 μm .

15

124. The method of any one of claims 112 to 123, wherein said gel and/or gel-forming substance is bioresorbable.

125. The method of claim 124, wherein said gel and/or gel-forming substance
20 comprises a biologically-based polymer(s) selected from the group consisting of collagen, fibrin, hyaluronan, chitosan and mixtures thereof.

126. The method of claim 124, wherein said gel and/or gel-forming substance
comprises a synthetic polymer(s) selected from the group consisting of
25 photopolymerizable end-capped block copolymers of poly(ethylene oxide) and an α -hydroxy acid.

127. The method of claim 124, wherein said gel and/or gel-forming substance
comprises a mixture of a biologically-based polymer(s) and a synthetic polymer(s),
30 wherein said biologically-based polymer(s) is/are selected from the group consisting of collagen, fibrin, hyaluronan, chitosan and mixtures thereof, and said synthetic polymer(s) is/are selected from the group consisting of photopolymerizable end-capped block copolymers of poly(ethylene oxide) and an α -hydroxy acid.

128. The method of any one of claims 124 to 127, wherein said gel and/or gel-forming substance is further comprised of a beneficial agent(s) selected from the group consisting of growth factors, glycosaminoglycans and hydrophilic compounds.

5 129. The method of any one of claims 124 to 128, wherein said gel and/or gel-forming substance includes an adhesive material(s).

130. The method of any one of claims 112 to 129, wherein an average of between about 3 and 500 cells and/or progenitor cells are associated with each of said beads or
10 particles.

131. The method of any one of claims 112 to 130, wherein said cells and/or progenitor cells are chondrocytes, embryonic stem cells and/or bone marrow stromal cells.

15

132. The method of any one of claims 112 to 130, wherein said cells and/or progenitor cells are fibroblast and/or progenitor cells thereof.

133. The method of any one of claims 112 to 130, wherein said cells and/or
20 progenitor cells are adipocytes and/or progenitor cells thereof.

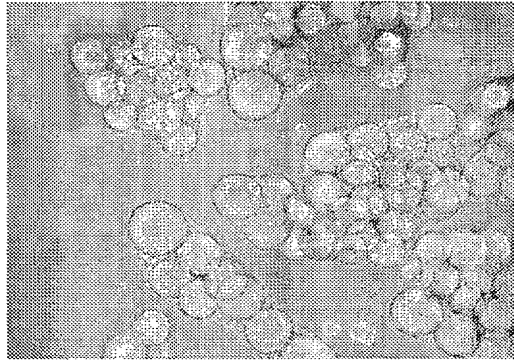
134. The method of any one of claims 112 to 130, wherein said cells and/or progenitor cells are osteoblasts and/or progenitor cells thereof.

25 135. The method of any one of claims 112 to 130, wherein said cells and/or progenitor cells are a mixture of cell types and/or progenitor cell types.

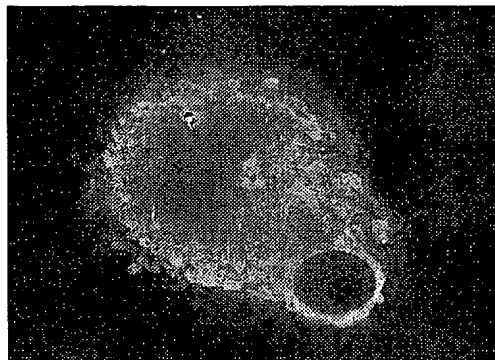
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1/4

Figure 1

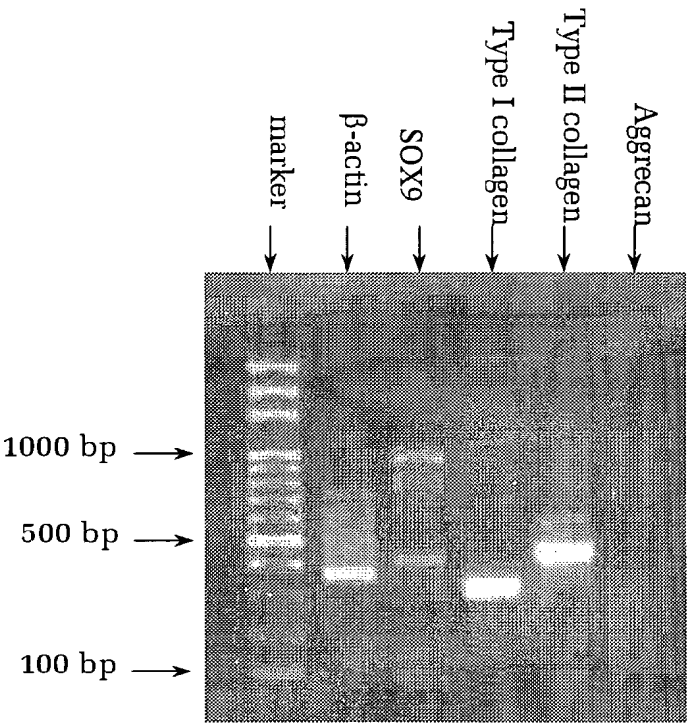


A



B

Figure 2



3/4

Figure 3

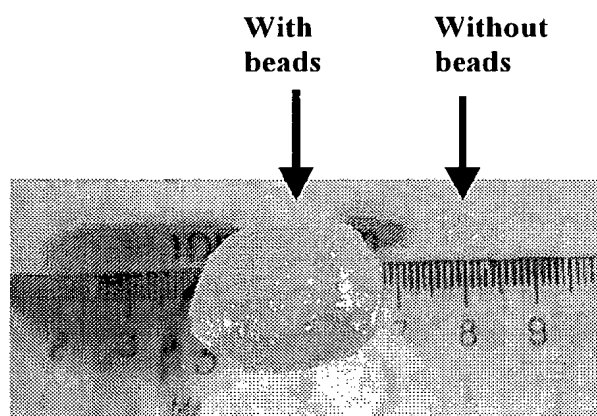
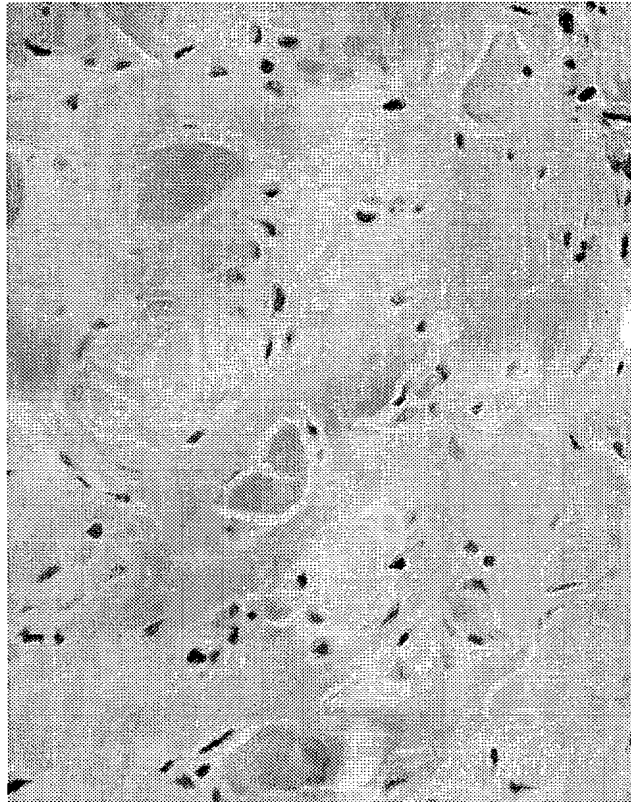


Figure 4



INTERNATIONAL SEARCH REPORT

International application No.

PCT/AU02/00106

A. CLASSIFICATION OF SUBJECT MATTER				
Int. Cl. ⁷ : A61K 35/32, 35/28, 35/48, 9/16; A61P 19/00				
According to International Patent Classification (IPC) or to both national classification and IPC				
B. FIELDS SEARCHED				
Minimum documentation searched (classification system followed by classification symbols)				
IPC A61K 35/32, 35/28, 35/48, 9/16				
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched				
AU: IPC AS ABOVE				
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)				
DERWENT: chondrocyte, stem cell, bone marrow, stromal cell, fibroblast, adipocyte, osteoblast, bead, particle				
C. DOCUMENTS CONSIDERED TO BE RELEVANT				
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.		
X	WO 99/15637 A (V.I. TECHNOLOGIES INC.), 1 April 1999 Page 13 lines 6-15, page 15 lines 3-18, page 16 lines 10-27; claims 23, 24, 41-43, 47 and 48	1-135		
P,X	WO 01/35932 A (THE REGENTS OF THE UNIVERSITY OF MICHIGAN), 25 May 2001 Page 6 lines 22-29, page 10 lines 13-25, page 20 lines 26-31, page 24 line 30- page 25 line 5; claims 1, 18, 38-39, 58 and 101	1-135		
P,X	US 2001/0014475 A (FRONDOZA et al.), 16 August 2001 Abstract, page 3, right col.; claims 1, 2, 4, 6, 9, 11, 21, 23, 48 and 50	1-135		
<input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C <input checked="" type="checkbox"/> See patent family annex				
<table border="0"> <tr> <td style="vertical-align: top;"> <p>* Special categories of cited documents:</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier application or patent but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> </td> <td style="vertical-align: top;"> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</p> <p>"&" document member of the same patent family</p> </td> </tr> </table>			<p>* Special categories of cited documents:</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier application or patent but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p>	<p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</p> <p>"&" document member of the same patent family</p>
<p>* Special categories of cited documents:</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier application or patent but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p>	<p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</p> <p>"&" document member of the same patent family</p>			
Date of the actual completion of the international search		Date of mailing of the international search report		
26 March 2002		- 3 APR 2002		
Name and mailing address of the ISA/AU		Authorized officer		
AUSTRALIAN PATENT OFFICE PO BOX 200, WODEN ACT 2606, AUSTRALIA E-mail address: pct@ipaustalia.gov.au Facsimile No. (02) 6285 3929		S. Chew Telephone No : (02) 6283 2248		

INTERNATIONAL SEARCH REPORT

International application No.

PCT/AU02/00106

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 00/56251 A (CHONDROS INC.), 28 September 2000 Whole document	1-135
A	WO 99/11196 A (POINT BIOMEDICAL CORP.), 11 March 1999 Whole document	1-135
A	WO 93/15722 A (SYNTEX (USA) INC.), 19 August 1993 Whole document	1-135

INTERNATIONAL SEARCH REPORT
Information on patent family members

International application No.
PCT/AU02/00106

This Annex lists the known "A" publication level patent family members relating to the patent documents cited in the above-mentioned international search report. The Australian Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

Patent Document Cited in Search Report				Patent Family Member			
WO	9915637	AU	94815/98	CA	2304384	EP	1015570
		US	6150505	US	2002006649		
WO	200135932	AU	200116228				
US	2001014475	US	2002012705	AU	200036267	EP	1171057
		WO	200056251	US	2001051834		